Unless otherwise noted, comments regarding the applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than January 27, 2010.

A. Federal Reserve Bank of Atlanta (Steve Foley, Vice President) 1000 Peachtree Street, N.E., Atlanta, Georgia 30309:
1. Florida Shores Bancorp, Inc., Smith Associates Bank Fund Management LLC, and Smith Associates Florida Banking Fund LLC, all of Pompano Beach, Florida; to collectively acquire at least 60 percent of the voting shares of Coastal Bancorporation, Inc., and thereby indirectly acquire voting shares of Coastal Bank, both of Merritt Island, Florida, and engage in operating a savings association, pursuant to section 225.28(b)(4)(ii) of Regulation Y. Comments regarding this application must be received by February 8, 2010.

B. Federal Reserve Bank of Dallas (E. Ann Worthy, Vice President) 2200 North Pearl Street, Dallas, Texas 75201–2272:
1. American Bank Holding Corporation, Corpus Christi, Texas; to engage de novo through its subsidiary, American Capital Solutions Group, Inc., Corpus Christi, Texas, in financial and investment advisory activities, pursuant to section 225.28(b)(6)(iii) of Regulation Y.


Robert deV. Frierson,
Deputy Secretary of the Board.
[FR Doc. 2010–362 Filed 1–11–10; 8:45 am]
BILLING CODE 6210–01–S

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

Sunshine Act; Notice of Meeting

TIME AND DATE: 9 a.m. (Eastern Time), January 19, 2010.
PLACE: 4th Floor Conference Room, 1250 H Street, NW., Washington, DC 20005.
STATUS: Parts will be open to the public and parts closed to the public.
MATTERS TO BE CONSIDERED: Parts Open to the Public:
1. Approval of the minutes of the November 16, 2009 Board member meeting.
2. Thrift Savings Plan activity report by the Executive Director.
   c. Legislative Report.
   d. Website Re-Design Update.
4. IT Modernization Plan Update.

Parts Closed to the Public
7. Confidential Financial Information.

CONTACT PERSON FOR MORE INFORMATION:
Thomas J. Trabucco, Director, Office of External Affairs, (202) 942–1640.
Thomas K. Emswiler,
Secretary, Federal Retirement Thrift Investment Board.
[FR Doc. 2010–441 Filed 1–8–10; 11:15 am]
BILLING CODE 6760–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Decision To Evaluate a Petition To Designate a Class of Employees for the General Electric Company, Evendale, OH, To Be Included in the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HHS gives notice as required by 42 CFR 83.12(e) of a decision to evaluate a petition to designate a class of employees for the General Electric Company, Evendale, Ohio, to be included in the Special Exposure Cohort under the Energy Employees Occupational Illness Compensation Program Act of 2000. The initial proposed definition for the class being evaluated, subject to revision as warranted by the evaluation, is as follows:

Facility: General Electric Company.
Location: Evendale, Ohio.
Job Titles and/or Job Duties: All employees of the Department of Energy, its predecessor agencies, and their contractors and subcontractors.

FOR FURTHER INFORMATION CONTACT:
Stuart L. Hinnefeld, Interim Director, Office of Compensation Analysis and Support, National Institute for Occupational Safety and Health (NIOSH), 4676 Columbia Parkway, MS C–46, Cincinnati, OH 45226, Telephone 513–533–6800 (this is not a toll-free number). Information requests can also be submitted by e-mail to OCASECDC.GOV.

John Howard,
Director, National Institute for Occupational Safety and Health.
[FR Doc. 2010–312 Filed 1–11–10; 8:45 am]
BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with the requirement for the opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Pub. L. 104–13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, e-mail paperwork@hrsa.gov or call the HRSA Reports Clearance Officer at (301) 443–1129.

Comments are invited on: (a) The proposal to continue collection of information for the proper performance of the functions of the Agency; (b) the accuracy of the Agency’s estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.


HRSA’s HIV/AIDS Bureau (HAB) administers Part A of Title XXVI of the Public Health Service Act as amended by Congress in October 2009 (Ryan White HIV/AIDS Treatment Extension Act of 2009). Part A provides emergency relief for areas with substantial need for HIV/AIDS care and support services that are most severely affected by the HIV/AIDS epidemic, including eligible metropolitan areas (EMA) and...
Transitional Grant Areas (TGAs). As a component of Part A (previously Title I), the purpose of the Minority AIDS Initiative (MAI) Supplement is to improve access to high quality HIV care services and health outcomes for individuals in disproportionately impacted communities of color who are living with HIV disease, including African-Americans, Latinos, Native Americans, Alaska Natives, Asian Americans, Native Hawaiians and Pacific Islanders (Section 2693(b)(2)(A) of the Public Health Service (PHS) Act).

Since the purpose of the Part A MAI is to expand access to medical, health, and social support services for disproportionately impacted racial/ethnic minority populations living with HIV/AIDS, who are not yet in care, it is important that HRSA is able to report on minorities served by the Part A MAI.

The Part A MAI Report is a data collection instrument in which grantees report on the number and characteristics of clients served and services provided. The Part A MAI Report, first approved for use in March 2006, is designed to collect performance data from Part A Grantees that will not change, and it has two parts: (1) A Web-based data entry application that collects standardized quantitative and qualitative information, and (2) an accompanying narrative report.

Grantees submit two Part A MAI Reports annually: Part A MAI Plan (Plan) and the Part A MAI Year-End Annual Report (Annual Report). The Plan and Annual Report components of the report are linked to minimize the reporting burden, and include drop-down menu responses, fields for reporting budget, expenditure and aggregated client level data, and open-ended responses for describing client or service-level outcomes. Together the Plan and Annual Report components collect information from grantees on MAI-funded services, expenditure patterns, the number and demographics of clients served, and client-level outcomes.

The MAI Plan Narrative that accompanies the Plan Web-forms provides (1) an explanation of the data submitted in the Plan Web forms; (2) a summary of the Plan, including the plan and timeline for disbursing funds, monitoring service delivery, and implementing any service-related capacity development or technical assistance activities; and (3) the plan and timeline for documenting client-level outcome measures. In addition, if the EMA/TGA revised any planned services, allocation amounts or target communities after their grant application was submitted, the changes must be highlighted and explained. The accompanying MAI Annual Report narrative describes (1) progress towards achieving specific goals and objectives identified in the Grantee’s approved MAI Plan for that fiscal year and in linking MAI services/activities to Part A and other Ryan White HIV/AIDS Program services; (2) achievements in relation to client-level health outcomes; (3) summary of challenges or barriers at the provider or grantee levels, the strategies and/or action steps implemented to address them, and lessons learned; and (4) discussion of MAI technical assistance needs identified by the EMA/TGA.

This information is needed to monitor and assess: (1) Changes in the type and amount of HIV/AIDS health care and related services being provided to each disproportionately impacted community of color; (2) the aggregate number of persons receiving HIV/AIDS services within each racial and ethnic community; and (3) the impact of Part A MAI-funded services in terms of client-level and service-level health outcomes. The information also is used to plan new technical assistance and capacity development activities and inform the HRSA policy and program management functions. The data provided to HRSA does not contain individual or personally identifiable information.

The annual estimated response burden for grantees is as follows:

<table>
<thead>
<tr>
<th>Form</th>
<th>Estimated number of respondents</th>
<th>Responses per respondent</th>
<th>Total responses</th>
<th>Hours per response</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Part A MAI Report</td>
<td></td>
<td></td>
<td>56</td>
<td>2</td>
<td>112</td>
</tr>
</tbody>
</table>

Note: Data collection system enhancements have resulted in a shortened response burden (from 6 to 5 total hours per response) for respondents since the previous OMB approval request.

E-mail comments to paperwork@hrsa.gov or mail the HRSA Reports Clearance Officer, Room 10–33, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857. Written comments should be received within 60 days of this notice.


Sahira Rafiullah,
Deputy Director, Division of Policy Review and Coordination.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0305]

Jason Vale; Denial of Hearing; Final Debarment Order

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is denying Jason Vale’s request for a hearing and is issuing an order under the Federal Food, Drug, and Cosmetic Act (the act) permanently debarring Mr. Vale from providing services in any capacity to a person that has an approved or pending drug product application. FDA bases this order on a finding that Mr. Vale was convicted of a felony under Federal law for conduct relating to the regulation of a drug product under the act. Mr. Vale has failed to file with the agency information and analyses sufficient to create a basis for a hearing concerning this action.

DATES: The order is effective January 12, 2010.

ADDRESSES: Submit applications for termination of debarment to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.


SUPPLEMENTARY INFORMATION: